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Claims

What is claimed is:

1.A compound represented by formula (1):

$$A - (CH_2)_m - X - (CH_2)_n$$
OH

wherein A is an optionally substituted phenyl or aromatic heterocyclic group which has 1 to 4 substituents selected from the group consisiting of a halogen atom, a hydroxyl group, an amino group, a nitro group, a cyano group, an alkyl group having 1 to 4 carbons, an alkoxy group having 1 to 4 carbons, an aminoalkyl group having 1 to 4 carbons, an acyl group having 1 to 4 carbons, an acylamino group having 1 to 4 carbons, an alkylthio group having 1 to 4 carbons, a perfluoroalkyl group having 1 to 4 carbons, a perfluoroalkyl group having 1 to 4 carbons, a perfluoroalkoxy group having 1 to 4 carbons, a carboxyl group, an alkoxycarbonyl group having 1 to 4 carbons, a phenyl group, an aromatic heterocyclic group and a heterocyclic group, said heterocyclic group being optionally substituted with an alkyl group having 1 to 4 carbons, a benzyl group, or a pyridylmethyl group;

m is an integer of 0 to 4;

n is an integer of 1 to 4;

X is a moiety having a structure selected from those illustrated in formula (2)

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— R¹ and R² are independently H or an optionally substituted alkyl group having 1 to 4 carbons; or a pharmaceutically acceptable salt thereof.

2.1.A compound of formula (1) according to claim 1 selected from the group consisting of

N-[4-(2-Hydroxycarbamoylvinyl)benzyl]-4-pyrrolidin-1-ylbenzamide, and 4-Dimethylamino-N-[4-(2-hydroxycarbamoylvinyl)benzyl]benzamide; or a pharmaceutically acceptable salt thereof.

- 3. 2. A pharmaceutical composition comprising a compound of formula (1) according to claim 1 in combination with a pharmaceutically acceptable excipient or diluent.
- 4. 3. Use of a compound according to claim 1 for the preparation of a medicament having histone deacetylase (HDAC) inhibitory activity.
 - 5. Use of a compound according to claim 4-as an inhibitor of cell-proliferation.
 - 6. Use of a compound according to claim 4 as an antitumor agent.